



REVOLUTIONARY SMART SELF ASSEMBLING  
CRYSTALLINE COLLOIDAL ARRAY MATERIALS FOR  
CONTROLLING IR LIGHT PROPAGATION

FINAL PROGRESS REPORT

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DECEMBER 13, 2000

U.S. ARMY RESEARCH OFFICE

GRANT NUMBER DAAG-55-98-1-0025

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# TECHNICAL PROGRESS REPORT

## A. STATEMENT OF PROBLEM

The work here developed new materials and fabrication methodologies to control IR light propagation.

1. We were to develop new syntheses for monodisperse polystyrene, silica, and PMMA colloids
2. Develop synthetic methods to produce colloidal particles containing quantum dots of CdS, CdTe, etc
3. Develop IR diffracting polymerized crystalline colloidal arrays (PCCA)
4. Develop methods to fabricate PCCA which diffract light in defined wavelengths and bandwidths
5. Fabricate PCCA which are thermally and chemically tunable in the 1-10  $\mu\text{m}$  spectral region
6. Demonstrate thermal tuning and electronic control of PCCA diffraction
7. Develop synthesis of highly charged, monodisperse titania dioxide colloids
8. Fabricate and characterize photonic bandgap crystals

## B. SUMMARY OF IMPORTANT RESULTS

1. We developed new methods to synthesize highly charged, monodisperse large polystyrene colloidal particles as reported in enclosed publication 1.
2. We developed novel methods to synthesize monodisperse silica colloids in microemulsions as shown in the synthetic scheme of Fig. 1, and the TEM showing examples of these colloidal particles in Fig. 2. We also developed synthetic methods to form silica particles which contain Ag, or Au quantum dots as shown in Figs. 3 and 4. We functionalized these silica composite particles to make them highly charged, such they self assembled into CCA as shown in Figs. 5 and 6. In addition, we developed methods to form polystyrene colloids which contained Ag quantum dots which also self assembled into CCA (Fig. 7 and 8).

We also developed methods using microemulsions to synthesize large silica particles containing CdS quantum dots (Fig. 9).

3. We developed a number methods to produce PCCA which diffract in the IR. In one method we hydrolyzed the acrylamide PCCA to form carboxylates. At neutral pH the charged carboxylates caused the PCCA to swell due to formation of a Donnan potential. Fig. 10 shows the resultant diffraction shift of a PCCA into the IR as the time for hydrolysis increases. This hydrolysis also creates a PCCA, which operates as a pH and ionic strength sensor. Our publication 2 is a careful study of the physics of the pH and ionic strength swelling, and its utility for sensing pH and ionic strength.

We also developed a chemically reversible disulfide cross linking approach (Fig. 11) and a hydrolysable crosslinking approach (Fig. 12 – 15) to swell PCCA into the IR.

4. The methods discussed above allow us to sensitively control the wavelength of diffraction of the PCCA by carefully adjusting the pH or ionic strength or by decreasing the PCCA crosslinking. We developed methods to control the diffraction bandwidth by altering the scattering power of the colloidal particles by changing their size or refractive index. We also attempted to swell colloidal particles already polymerized into the PCCA. Preliminary attempts to swell polystyrene particles by adding styrene to the PCCA solution indicates success. Further work will be necessary to speed up this process in order to make it practical.

5. We demonstrated that we could achieve thermal tuning by utilizing PCCA synthesized by using hydrogels of N-isopropyl acrylamide.

We began working on electronic control of the PCCA diffraction by utilizing a PCCA which contains an 18-crown-6 crown ether group. This molecular recognition agent binds  $Pb^{2+}$ , which causes the PCCA to swell and red shifts the diffraction. We attempted electronic control of the PCCA diffraction by utilizing the electrochemical oxidization of lead metal to  $Pb^{2+}$  which would then swell the PCCA into the IR. Preliminary data suggests that this method is viable for the control of PCCA diffraction wavelengths.

6. The most challenging part of the research program involved synthesis of highly charged, monodisperse titania spheres. Although the literature reports synthesis of large, relatively monodisperse titania spheres, we found that none of these recipes were reproducible and did not yield monodisperse colloidal particles. We developed a number of new approaches for synthesizing titania spheres of relatively uniform size, but when we tried to add surface charge the titania colloids suffered, what we think, was a Coulomb explosion where the particles disintegrated into their small primary particles. Only just recently, with funding from other grant support, have we finally succeeded in fabricating monodisperse titania colloidal particles which self assemble into CCA that Bragg diffract light. Fig. 16 shows TEM micrographs of these particles.

7. We reported earlier that colloidal arrays made from polystyrene colloids with a lattice spacing that diffracts  $\sim 1200$  nm light from the fcc 111 planes diffracts essentially all light of all polarizations at wavelengths below 500 nm. Clearly the refractive index contrast, and the fact that it is an fcc crystal will not allow this sample to be a 3-D photonic bandgap crystal. We have been carefully characterizing the crystal structure and diffraction properties and writing up these results for publication. We expect to submit this work by the end of December, 2000.

#### C. PUBLICATIONS

1. C. Reese, C. Guerrero, J. Weissman, K. Lee and S. A. Asher, "Synthesis of Highly Charged, Monodisperse Polystyrene Colloidal Particles for the Fabrication of Photonic Crystals", *Journal of Colloid and Interface Science*, **232**, 76-80 (2000)
2. K. Lee and S. A. Asher, "Photonic Crystal Chemical Sensors: pH and Ionic Strength", *Journal of American Chemical Society*, **122**, 9534-9537 (2000)

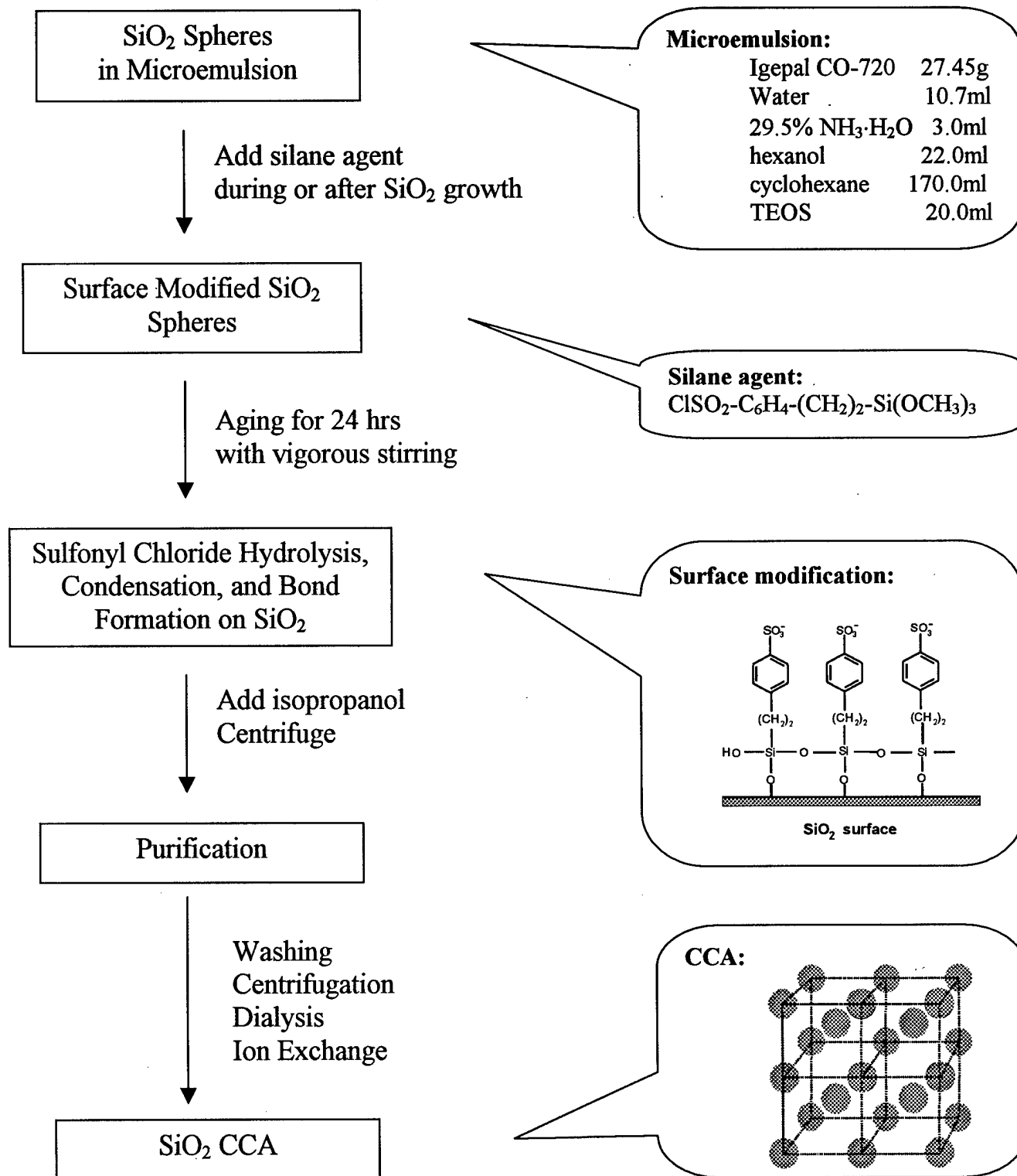
#### D. LIST OF SCIENTIFIC PERSONNEL EMPLOYED ON PROJECT

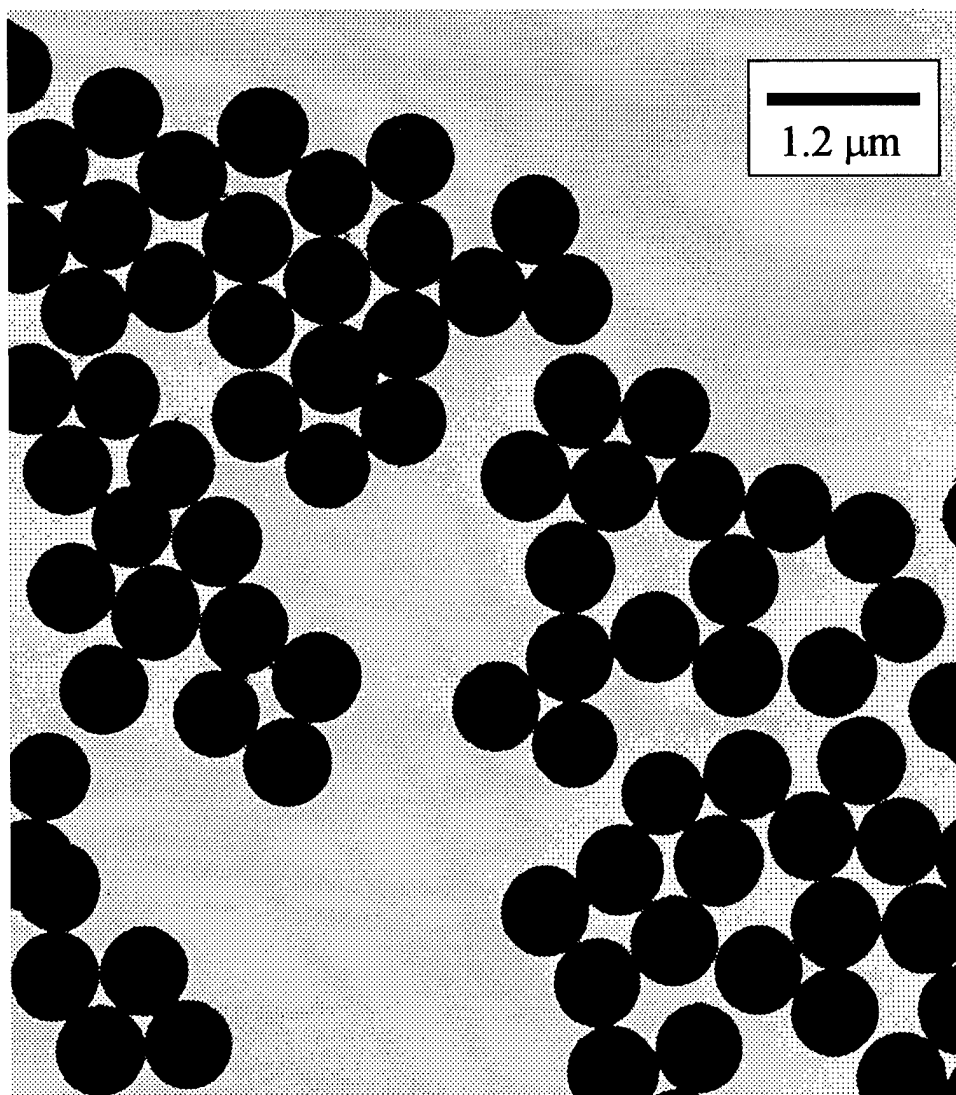
Sanford Asher	PI	12%
We Wang	Res. Associate	50%
Igor Lednev	Res. Associate	50%
Jeff Sicher	Machinist	4%
Douglas Hackworth	Graduate Student	75%
Jonathan Keim	Graduate Student	25%
Aleksandr Mikhonin	Graduate Student	25%
Vasil Pajcini	Graduate Student	25% Received PhD
Hua Zhang	Graduate Student	75% Received MS

#### E. REPORT ON INVENTIONS

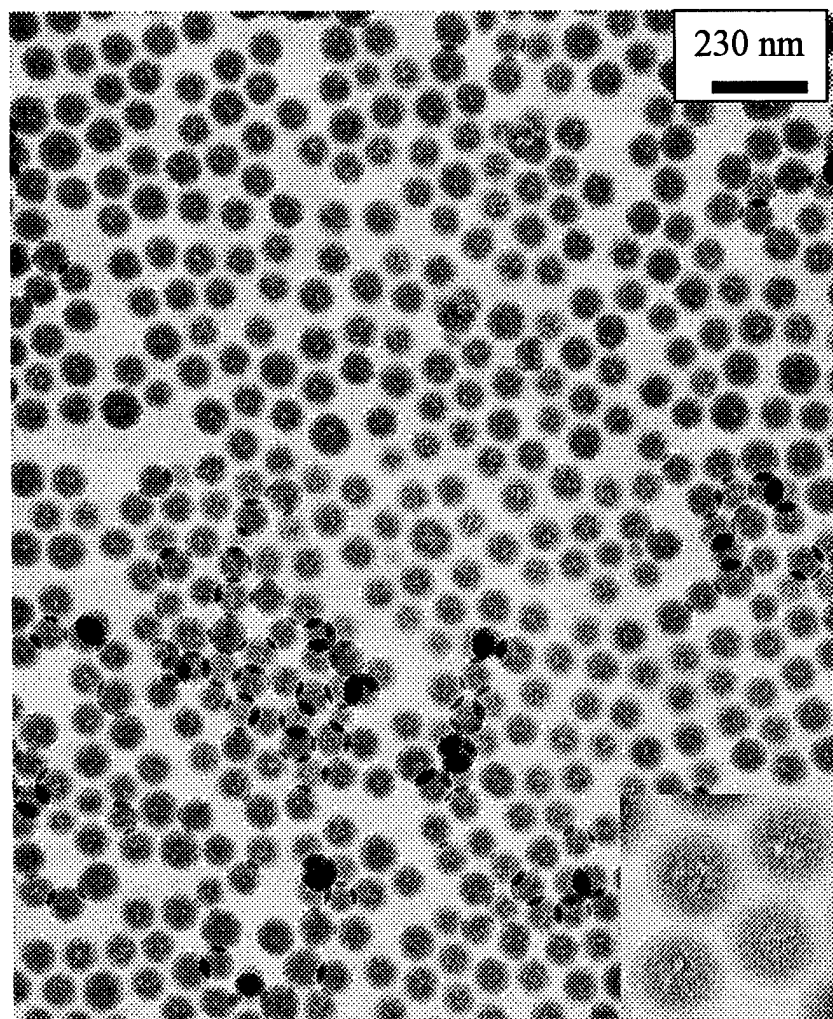
None, as yet.

**Figure 1. Synthesis of Highly Charged, Monodisperse SiO<sub>2</sub> Particles**



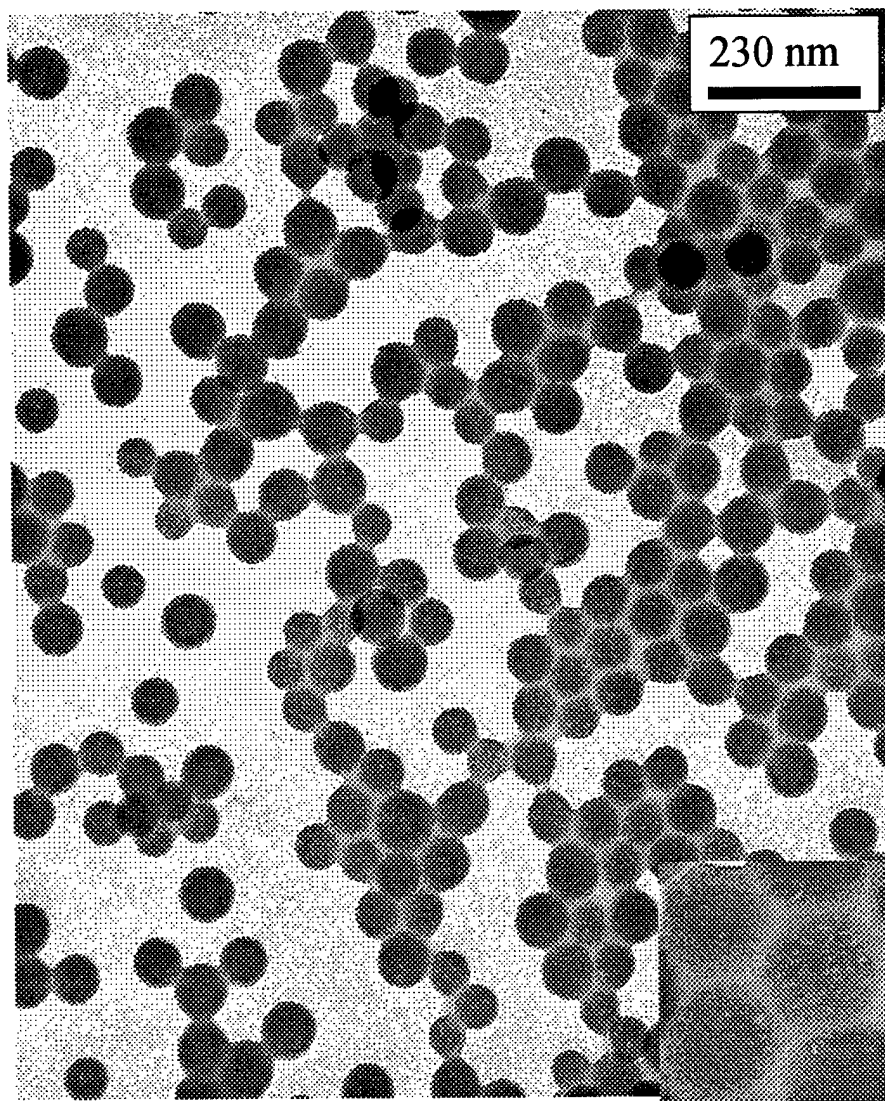


***Figure 2.*** TEM Picture of Surface Modified SiO<sub>2</sub> Particles

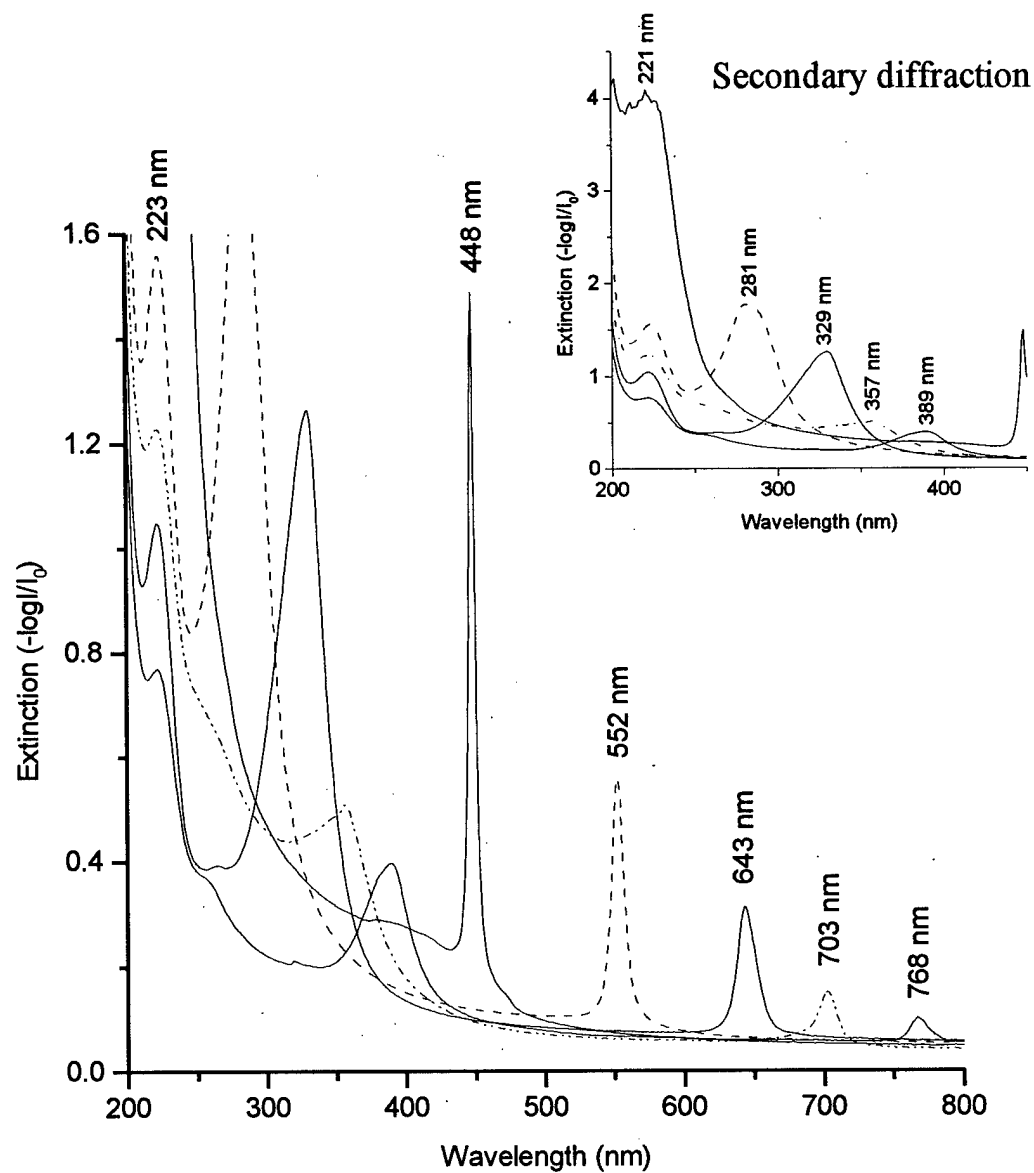


**Figure 3.** TEM picture of SiO<sub>2</sub>-Ag composite particles (Ag homogeneously distribution in SiO<sub>2</sub> spheres)

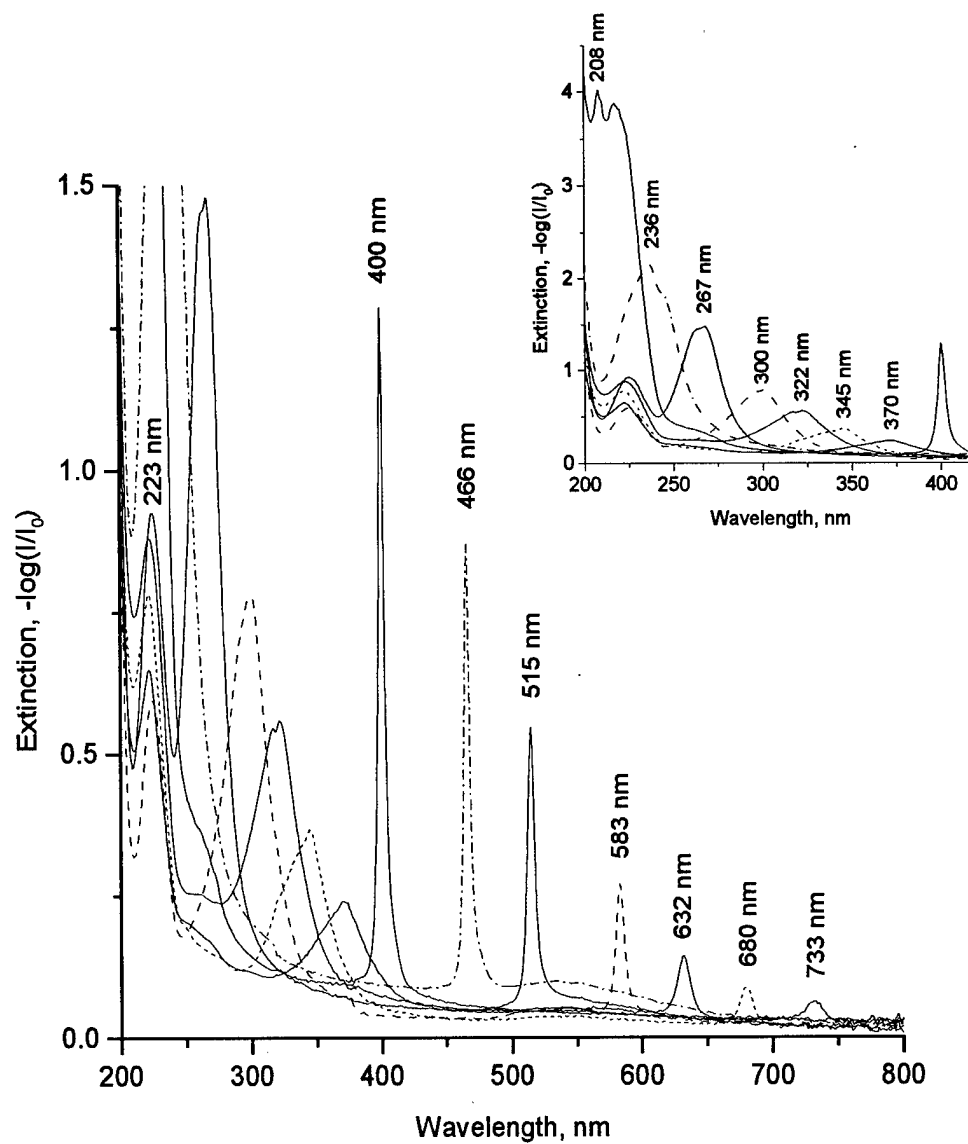




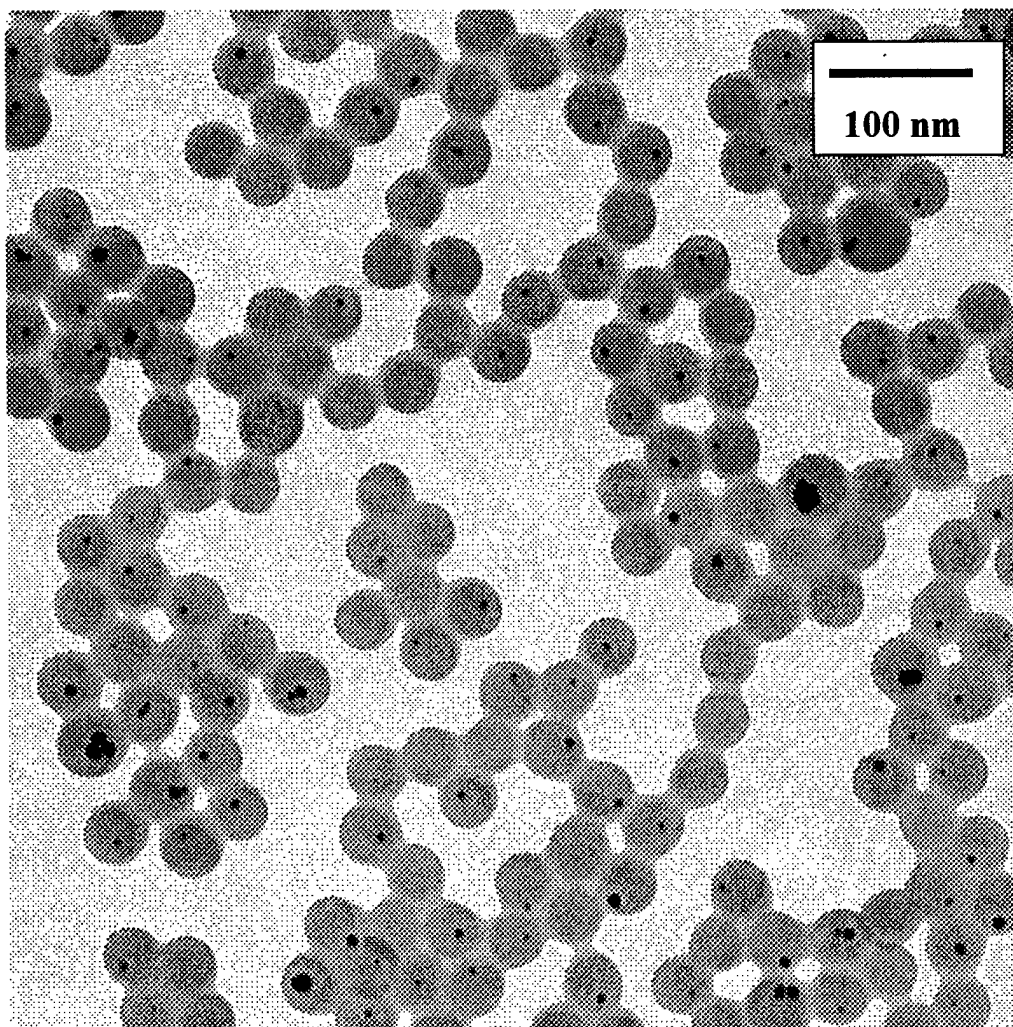
**Figure 4.** TEM picture of SiO<sub>2</sub>-Au composite spheres. The Au nanoparticles homogeneously distribute in the SiO<sub>2</sub> spheres.



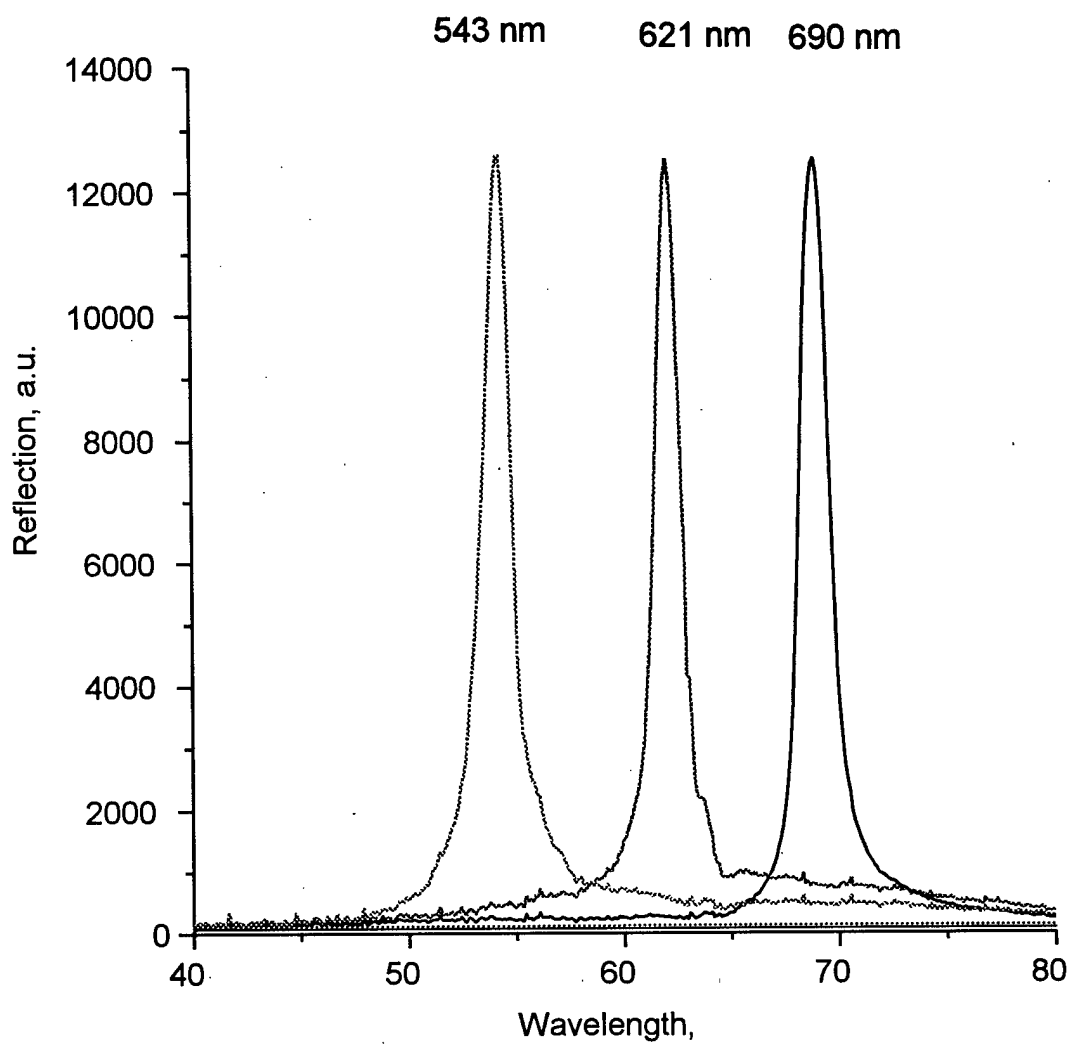
**Figure 5.** Extinction spectra of Ag@SiO<sub>2</sub> composite sphere CCA at different colloidal concentrations. The Ag@SiO<sub>2</sub> sphere diameter is 98±8 nm.



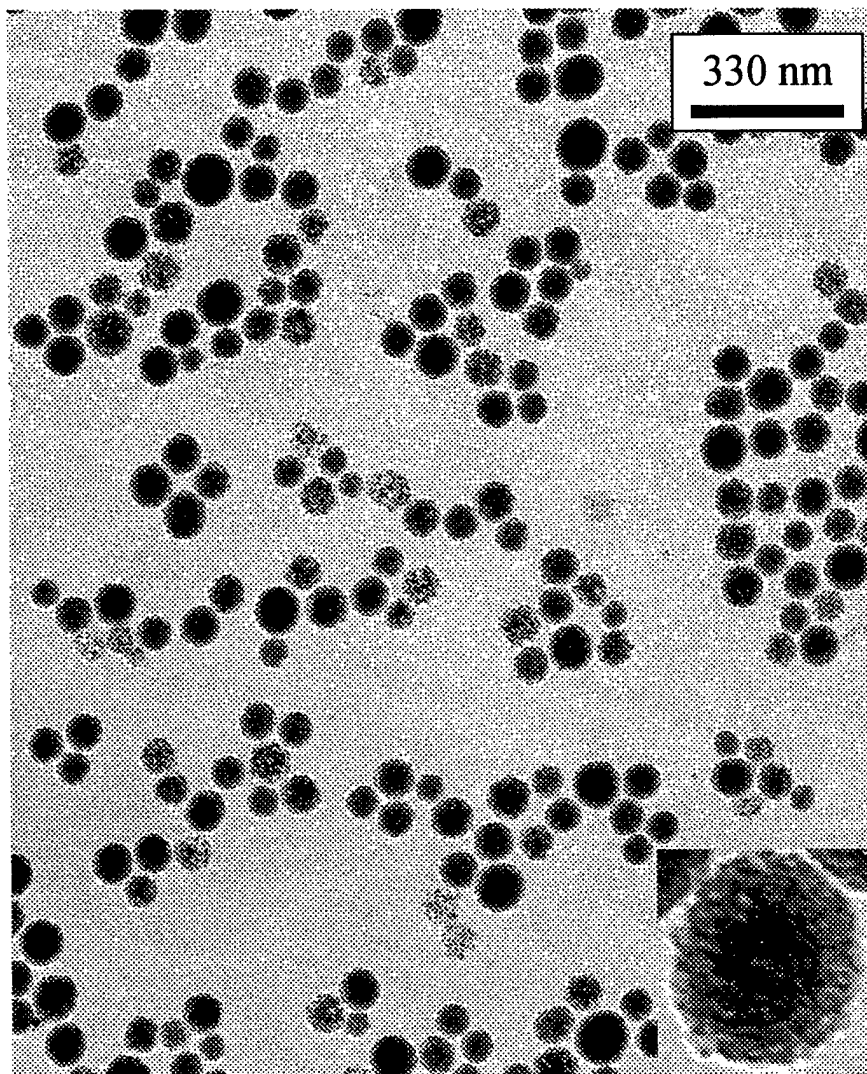
**Figure 6.** Extinction spectra of Au@SiO<sub>2</sub> composite sphere CCA at different sphere concentrations. The sphere diameter is  $110 \pm 10$  nm, and the thickness of the CCA is 0.125 mm.



*Figure 7.* TEM picture of polystyrene-silver shell-core particles.

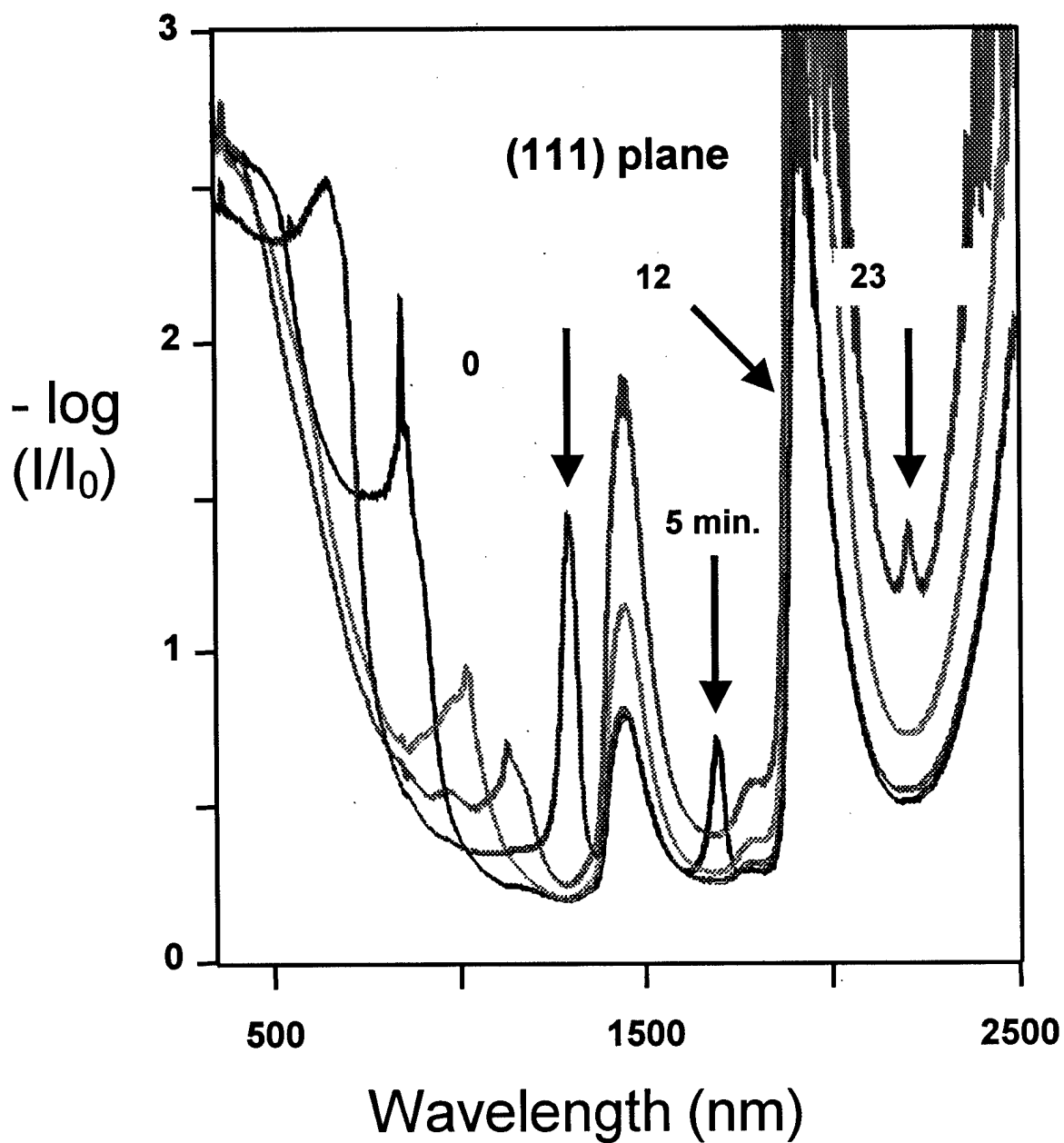


**Figure 8.** Back-scattering diffraction spectra of polystyrene-silver shell-core composite CCA at different particle concentrations.



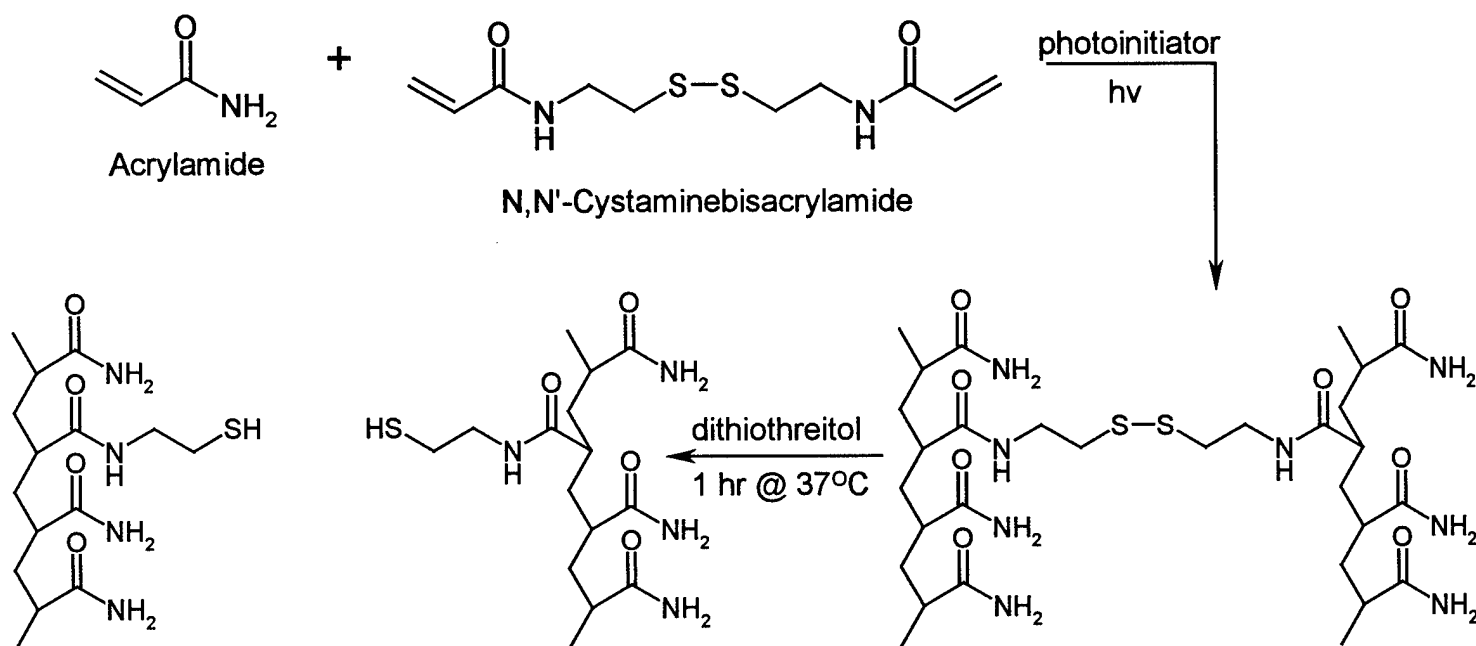
**Figure 9.** TEM picture of SiO<sub>2</sub>-CdS (surface) composite particles

# PCCA SWELLED INTO IR



**Figure 10.** PS Colloid 270 nm diameter, Acrylamide PCCA Hydrolyzed for 0, 5, 12, 23 minutes.

# Increasing PCCA Spacing and Decreasing Elastic Constant Chemically Variable Crosslinking



Hydrogel with cleaved crosslinks

Crosslinked hydrogel

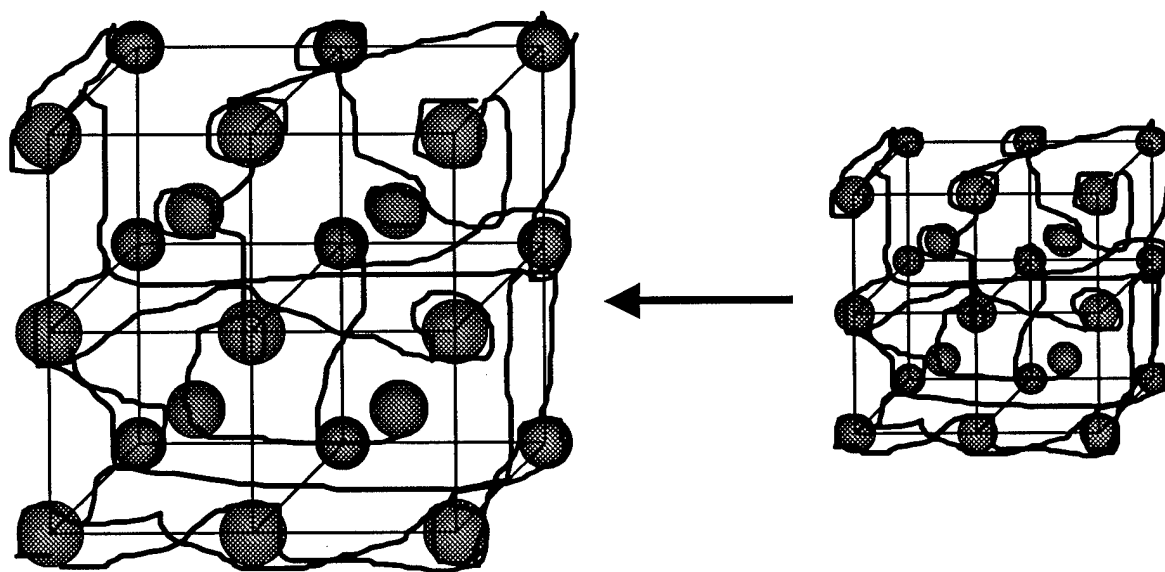
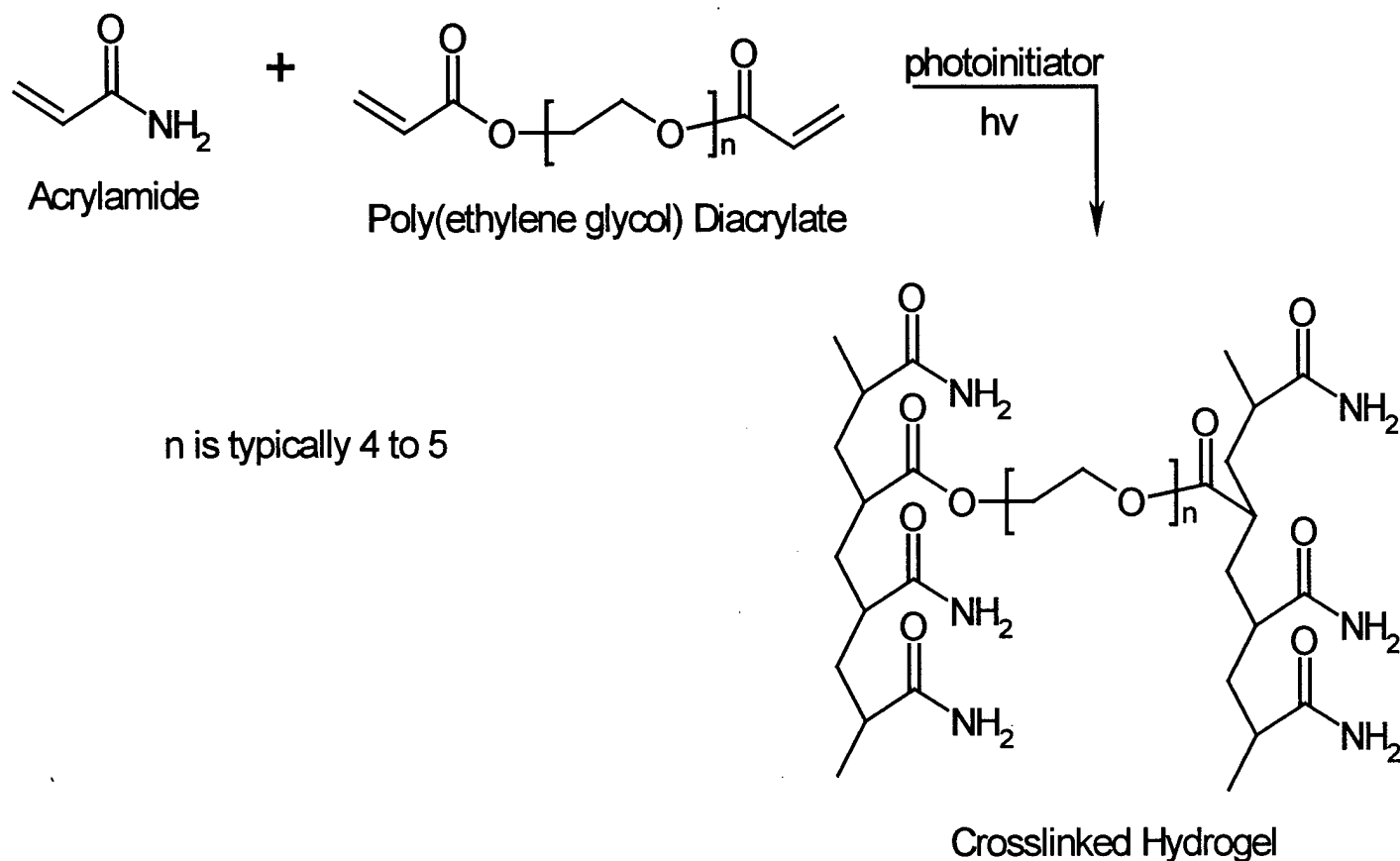


Figure 11



# Fabrication of IR Diffracting Hydrogels

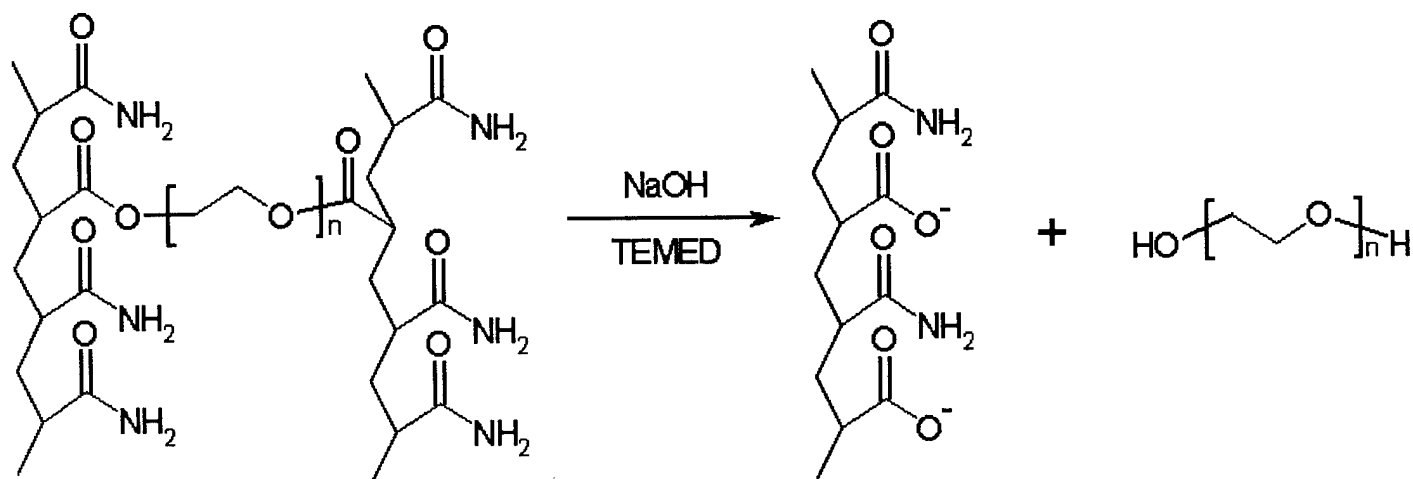
## Synthesis of PEG Crosslinked Hydrogels



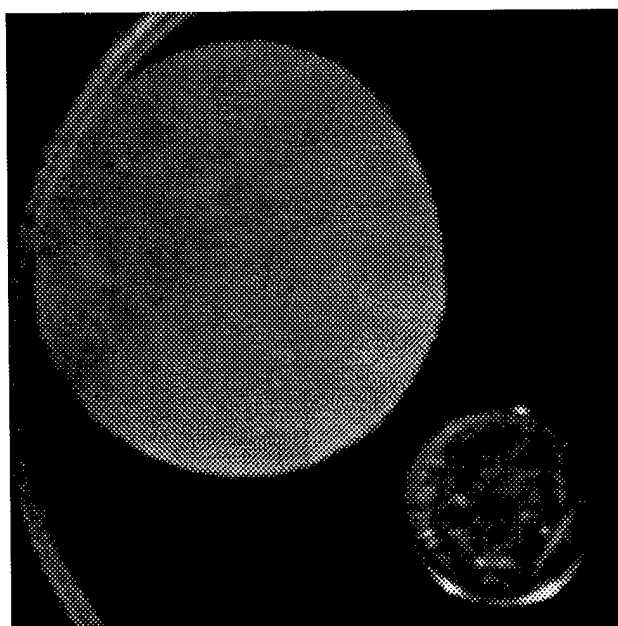
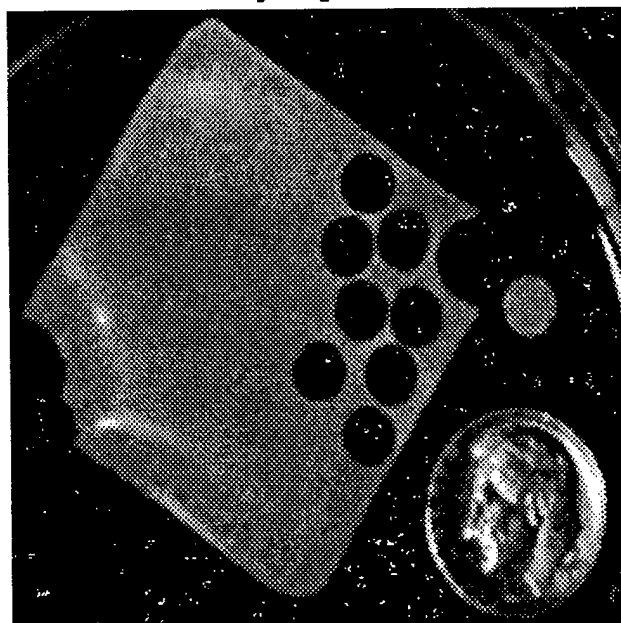
*Figure 12*

# Fabrication of IR Diffracting Hydrogels

## Hydrolysis of Crosslinks



Crosslinked Hydrogel



### Typical Recipe:

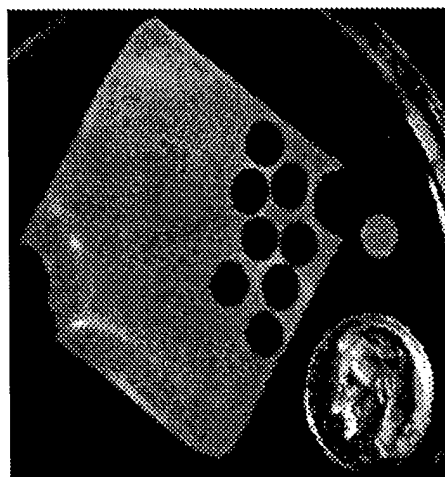
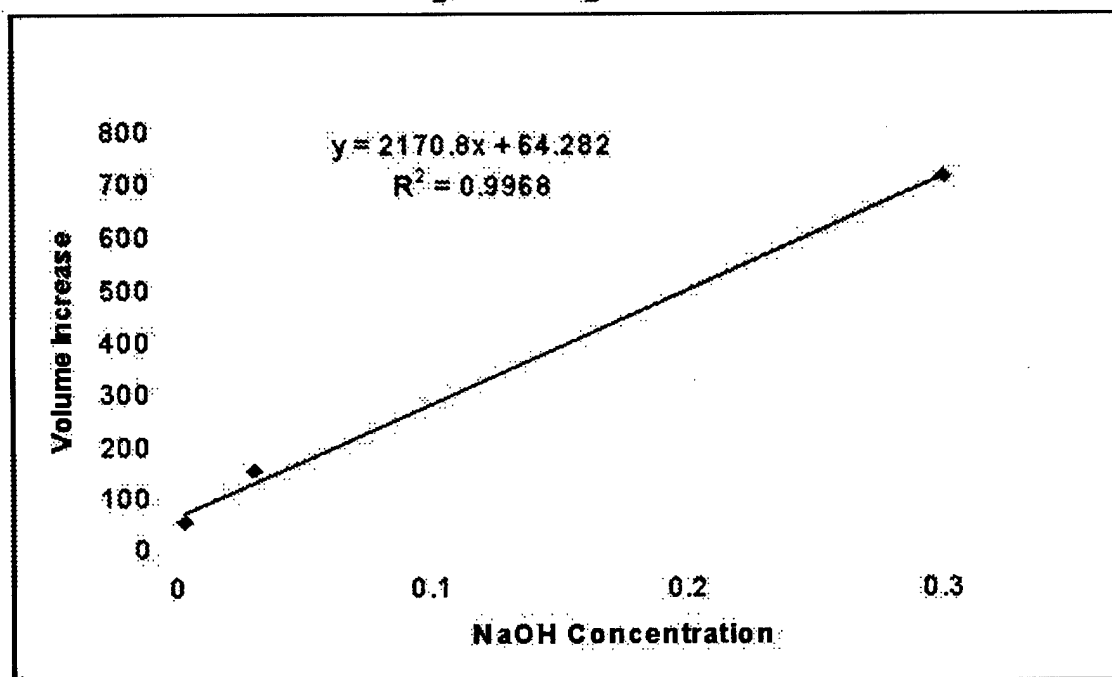
0.03 M NaOH

0.08% N,N,N',N'-tetramethylethylenediamine (TEMED)

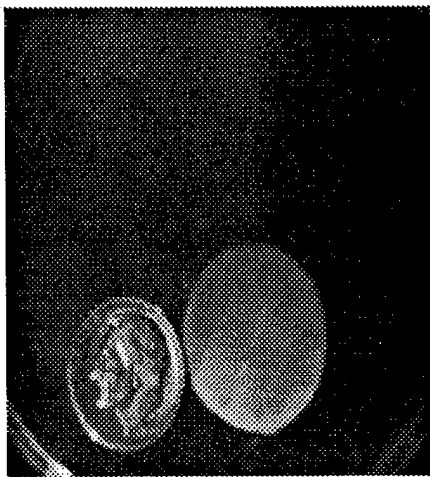
for 1 hour

*Figure 13*

# Gel Volume Increase With Changing NaOH Concentration After 1 Hour Of Hydrolysis



**Gel Before Hydrolysis**



**After Hydrolysis in 0.03 M NaOH**

$$\frac{V_f}{V_i} = 150$$



**After Hydrolysis in 0.3 M NaOH**

$$\frac{V_f}{V_i} = 10^3$$

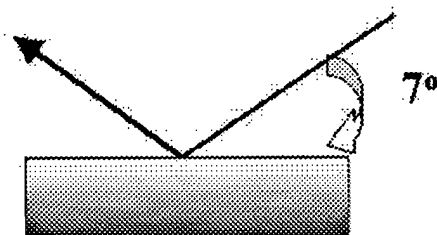
*Figure 14*

## Diffraction Still Present After Swelling

$$\lambda_{111}^i = 950 \text{ nm} \quad d_{111}^i = 354 \text{ nm}$$

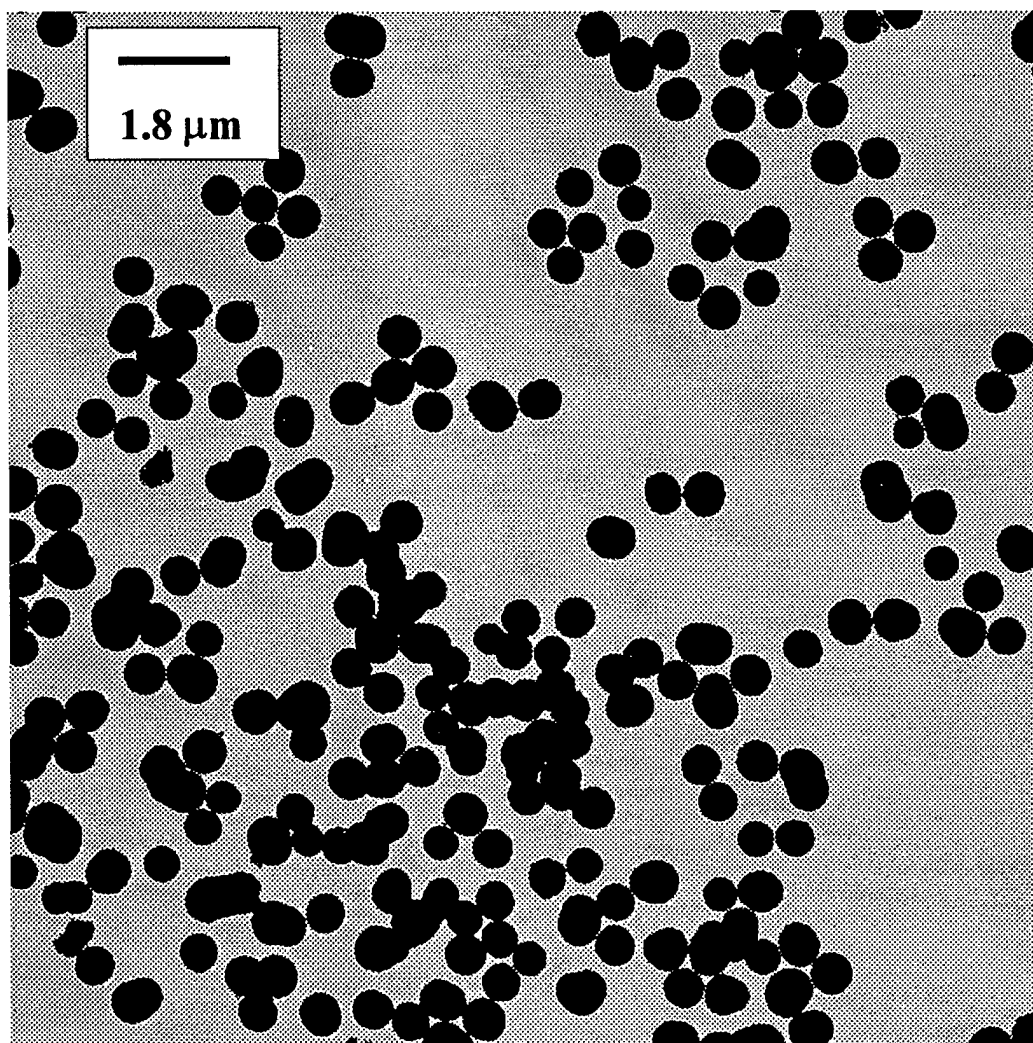
$$\frac{V_f}{V_i} = 150$$

$$\lambda_{111}^f = 5.035 \mu\text{m} \text{ (Normal incidence)}$$



$$\lambda_d^{7^\circ} = 608 \text{ nm}$$

Figure 15



*Figure 16.* TEM picture of monodisperse TiO<sub>2</sub> spheres.